CLAIM AMENDMENTS

1. - 39. (canceled)

40. (currently amended) A compound of formula A or formula B:

where

R¹, R², and R³ are independently, hydrogen, or optionally substituted lower alkyl, alkenyl, alkynyl, cycloalkyl, cycloalkyl(lower alkyl), optionally substitutedheterocycloalkyl, optionally substituted aryl, optionally substituted heteroaryl, optionally substituted aryl(lower alkyl), halo(lower alkyl), -CF₃, halogen, nitro, -CN, -OR⁹, -SR⁹, -NR⁹R¹⁰, -NR⁹(carboxy(lower alkyl)), -C(=O)R⁹, -C(=O)OR⁹, -C(=O)NR⁹R¹⁰, -OC(=O)R⁹, -SO₂R⁹, -OSO₂R⁹, -SO₂NR⁹R¹⁰, -NR⁹SO₂R¹⁰ or -NR⁹C(=O)R¹⁰, where R⁹ and R¹⁰ are independently, hydrogen, optionally substituted lower alkyl, lower alkyl-N(C₁₋₂ alkyl)₂, lower alkyl(optionally substituted heterocycloalkyl), alkenyl, alkynyl, optionally substituted cycloalkyl, cycloalkyl(lower alkyl), optionally substituted heterocycloalkyl(lower alkyl), optionally substituted aryl, optionally substituted aryloxy, heteroaryl, heteroaryl(lower alkyl), or R⁹ and R¹⁰ together are -(CH₂)₄₋₆- optionally interrupted by one O, S, NH, N-(aryl), N-(aryl(lower alkyl)), N-(carboxy(lower alkyl)) or N-(optionally substituted C₁₋₂ alkyl) group,

R⁴ and R⁵ are independently, hydrogen or lower alkyl optionally substituted lower alkyl, optionally substituted aryl, or optionally substituted aryl(lower-alkyl), or, together, are (CH₂)₂₋₄-,

R⁶ is hydrogen, optionally substituted lower alkyl, alkenyl, alkynyl, cycloalkyl, cycloalkyl(lower alkyl), optionally substituted heterocycloalkyl, optionally substituted aryl, optionally substituted heteroaryl, optionally substituted

heteroaryl(lower alkyl), -C(=O)R¹¹, -C(=O)OR¹¹, -C(=O)NR¹¹R¹², -SO₂R¹¹, or -SO₂NR¹¹R¹², where R ¹¹ and R¹² are independently, hydrogen, optionally substituted lower alkyl, alkenyl, alkynyl, cycloalkyl, cycloalkyl(lower alkyl), aryl, optionally substituted aryloxy, heteroaryl, heteroaryl(lower alkyl), or R¹¹ and R¹² together are -(CH₂)₄₋₆-,

or a pharmaceutically acceptable salt thereof, optionally in the form of a single stereoisomer or mixture of stereoisomers thereof.

- 41. (previously presented) The compound of claim 40, where said compound is a compound of Formula A or a pharmaceutically acceptable salt thereof, optionally in the form of a single stereoisomer or mixture of stereoisomers thereof.
- 42. (previously presented) The compound of claim 40, where said compound is a compound of Formula B or a pharmaceutically acceptable sait thereof, optionally in the form of a single stereoisomer or mixture of stereoisomers thereof.
- 43. (previously presented) The compound of claim 40, where R¹ is hydrogen, optionally substituted lower alkyl, optionally substituted heterocycloalkyl, optionally substituted aryl, optionally substituted heteroaryl, optionally substituted aryl(lower alkyl), halogen, -OR⁹, -NR⁹R¹⁰, -C(=O)OR⁹, -C(=O)NR⁹R¹⁰, -SO₂NR⁹R¹⁰, or -NR⁹C(=O)R¹⁰, where R⁹ and R¹⁰ are independently, hydrogen, optionally substituted lower alkyl, lower alkyl-N(C₁₋₂ alkyl)₂, lower alkyl(optionally substituted heterocycloalkyl), aryl(lower alkyl), optionally substituted aryl, heteroaryl, or heteroaryl(lower alkyl).
- 44. (previously presented) The compound of claim 43, where R¹ is optionally substituted lower alkyl, optionally substituted heterocycloalkyl, optionally substituted aryl, optionally substituted aryl(lower alkyl), halogen, -OR⁹, -NR⁹R¹⁰, -C(=O)OR⁹, -C(=O)NR⁹R¹⁰, -SO₂NR⁹R¹⁰, or -NR⁹C(=O)R¹⁰, where R⁹ and R¹⁰ are independently, hydrogen, optionally substituted lower alkyl, lower alkyl-N(C₁₋₂ alkyl)₂, lower alkyl(optionally substituted heterocycloalkyl), aryl(lower alkyl), optionally substituted aryl, heteroaryl, or heteroaryl(lower alkyl).
- 45. (previously presented) The compound of claim 40, where R² is hydrogen, optionally substituted lower alkyl, cycloalkyl, optionally substituted heterocycloalkyl, optionally substituted

aryl, optionally substituted heteroaryl, optionally substituted aryl(lower alkyl), halogen,-OR⁹, -NR⁹(carboxy(lower alkyl)), -C(=O)OR⁹, -C(=O)NR⁹R¹⁰, -SO₂NR⁹R¹⁰, or -NR⁹C(=O)R¹⁰, where R⁹ and R¹⁰ are independently, hydrogen, optionally substituted lower alkyl, lower alkyl-N(C₁₋₂ alkyl)₂, lower alkyl(optionally substituted heterocycloalkyl), optionally substituted cycloalkyl, cycloalkyl(lower alkyl), optionally substituted aryl, optionally substituted aryloxy, heteroaryl, heteroaryl(lower alkyl), or R⁹ and R¹⁰ together are -(CH₂)₄₋₆- optionally interrupted by one O, S, NH, N-(aryl), N-(aryl(lower alkyl)), N-(carboxy(lower alkyl)) or N-(optionally substituted C₁₋₂ alkyl) group.

46. (previously presented) The compound of claim 45, where R² is optionally substituted lower alkyl, cycloalkyl, optionally substituted heterocycloalkyl, optionally substituted aryl, optionally substituted heteroaryl, optionally substituted aryl(lower alkyl), halogen, -OR⁹, -NR⁹(carboxy(lower alkyl)), -C(=O)OR⁹, -C(=O)NR⁹R¹⁰, -SO₂NR⁹R¹⁰, or -NR⁹C(=O)R¹⁰, where R⁹ and R¹⁰ are independently, hydrogen, optionally substituted lower alkyl, lower alkylN(C₁₋₂ alkyl)₂, lower alkyl(optionally substituted heterocycloalkyl), optionally substituted cycloalkyl, cycloalkyl(lower alkyl), optionally substituted aryl, optionally substituted aryloxy, heteroaryl, heteroaryl(lower alkyl), or R⁹ and R¹⁰ together are -(CH₂)₄₋₆- optionally interrupted by one O, S, NH, N-(aryl), N-(aryl(lower alkyl)), N-(carboxy(lower alkyl)) or N-(optionally substituted C₁₋₂ alkyl) group.

47. (previously presented) The compound of claim 40, where R³ is hydrogen, optionally substituted lower alkyl, optionally substituted heterocycloalkyl, optionally substituted aryl, optionally substituted heteroaryl, optionally substituted aryl(lower alkyl), halo(lower alkyl), halogen, -OR⁹, -NR⁹R¹⁰, -C(=O)OR⁹, or -C(=O)NR⁹R¹⁰, where R⁹ and R¹⁰ are independently, hydrogen, optionally substituted lower alkyl, lower alkyl-N(C₁₋₂ alkyl)₂, lower alkyl(optionally substituted heterocycloalkyl), optionally substituted cycloalkyl, cycloalkyl(lower alkyl), optionally substituted aryloxy, heteroaryl, heteroaryl(lower alkyl), or R⁹ and R¹⁰ together are -(CH₂)₄₋₆- optionally interrupted by one O, S, NH, N-(aryl), N-(aryl(lower alkyl)), N-(carboxy(lower alkyl)) or N-(optionally substituted C₁₋₂ alkyl) group.

48. (previously presented) The compound of claim 47, where R³ is optionally substituted lower alkyl, optionally substituted heterocycloalkyl, optionally substituted aryl, optionally

substituted heteroaryl, optionally substituted aryl(lower alkyl), halo(lower alkyl), halogen, -OR⁹, -NR⁹R¹⁰, -C(=0)OR⁹, or -C(=0)NR⁹R¹⁰, where R⁹ and R¹⁰ are independently, hydrogen, optionally substituted lower alkyl, lower alkyl-N(C₁₋₂ alkyl)₂, lower alkyl(optionally substituted heterocycloalkyl), optionally substituted cycloalkyl, cycloalkyl(Iower alkyl), optionally substituted aryl, optionally substituted aryloxy, heteroaryl, heteroaryl (lower alkyl), or R9 and R10 together are -(CH₂)₄₋₆- optionally interrupted by one O, S, NH, N-(aryl), N-(aryl(lower alkyl)), N-(carboxy(lower alkyl)) or N-(optionally substituted C₁₋₂ alkyl) group.

49. (canceled)

The compound of claim 40, where R⁶ is hydrogen, optionally 50. (previously presented) substituted lower alkyl, alkenyl, cycloalkyl, cycloalkyl(lower alkyl), optionally substituted heterocycloalkyl, optionally substituted aryl, optionally substituted aryl(lower alkyl), optionally substituted heteroaryl, optionally substituted heteroaryl(lower alkyl), -C(=0)R¹¹, -C(=0)OR¹¹, -C(=O)NR¹¹R¹², -SO₂R¹¹, or -SO₂NR¹¹R¹², where R¹¹ and R¹² are independently, hydrogen, optionally substituted lower alkyl, cycloalkyl, cycloalkyl(lower alkyl), aryl heteroaryl, heteroaryl(lower alkyl), or R11 and R12 together are -(CH2)4-6-.

51. (previously presented) The compound of claim 40 that is a compound of formula Aa or formula Ba:

$$R^{1}$$
 R^{2}
 R^{2}
 R^{3}
 R^{4}
 R^{5}
 R^{13}
 R^{13}

where:

R¹, R², R³, R⁴, and R⁵ are as defined in claim 40,

R¹³ is hydrogen, optionally substituted lower alkyl, alkenyl, alkynyl, cycloalkyl, cycloalkyl(lower alkyl), heterocycloalkyl, optionally substituted aryl, optionally substituted aryl(lower alkyl), optionally substituted heteroaryl, optionally substituted heteroaryl(lower alkyl), halo(lower alkyl), -CF₃, halogen, nitro, -CN, -OR¹⁵, -SR¹⁵, -NR¹⁵R¹⁶, -C(=O)R¹⁵,

-C(=O)OR¹⁵, -C(=O)NR¹⁵R¹⁶, -OC(=O)R¹⁵, -SO₂R¹⁵, -SO₂NR¹⁵R¹⁶, -NR¹⁵SO₂R¹⁶ or -NR¹⁵C(=O)R¹⁶, where R¹⁵ and R¹⁶ are independently, hydrogen, optionally substituted lower alkyl, alkenyl, alkynyl, -CF₃, cycloalkyl, optionally substituted heterocycloalkyl, cycloalkyl(lower alkyl), optionally substituted aryloxy, optionally substituted heteroaryl, optionally substituted heteroaryl(lower alkyl), or, together, are -(CH₂)₄₋₆- optionally interrupted by one O, S, NH or N-(C₁₋₂ alkyl) group,

each R¹⁴ is independently selected from optionally substituted lower alkyl, optionally substituted aryl, optionally substituted heteroaryl, hydroxy, halogen, -CF₃, -OR¹⁷, -NR¹⁷R¹⁸, -C(=O)R¹⁸, -C(=O)OR¹⁸, -C(=O)NR¹⁷R¹⁸, where R¹⁷ and R¹⁸ are independently, hydrogen, lower alkyl, alkenyl, alkynyl, -CF₃, optionally substituted heterocycloalkyl, cycloalkyl, cycloalkyl(lower alkyl), optionally substituted aryl, optionally substituted aryloxy, heteroaryl, heteroaryl(lower alkyl), or, together, are-(CH₂)₄₋₆-, optionally interrupted by one O, S, NH or N-(C₁₋₂ alkyl) group, and

where n is an integer of 0 to 4,

or a pharmaceutically acceptable salt thereof, optionally in the form of a single stereoisomer or mixture of stereoisomers.

- 52. (previously presented) The compound of claim 51, where said compound is a compound of Formula Aa or a pharmaceutically acceptable salt thereof, optionally in the form of a single stereoisomer or mixture of stereoisomers.
- 53. (previously presented) The compound of claim 51, where said compound is a compound of Formula Ba or a pharmaceutically acceptable salt thereof, optionally in the form of a single stereoisomer or mixture of stereoisomers.
- 54. (previously presented) The compound of claim 51, where R¹³ is -OR¹⁵, and R¹⁵ is hydrogen, lower alkyl optionally substituted with -C(=0)OR¹⁹, where R¹⁹ is hydrogen or lower alkyl, alkenyl, alkynyl, -CF₃, cycloalkyl, optionally substituted heterocycloalkyl, cycloalkyl(lower alkyl), optionally substituted aryl, optionally substituted heteroaryl, or optionally substituted heteroaryl(lower alkyl).
- 55. (previously presented) The compound of claim 51, where R¹³ is hydrogen, optionally substituted lower alkyl, alkenyl, heterocycloalkyl, optionally substituted aryl, optionally

substituted aryl(lower alkyl), optionally substituted heteroaryl, optionally substituted heteroaryl(lower alkyl), halo(lower alkyl), -CF₃, halogen, nitro, -CN, -OR¹⁵, -SR¹⁵, -NR¹⁵R¹⁶, -C(=O)R¹⁵, -C(=O)R¹⁵, -C(=O)NR¹⁵R¹⁶, -OC(=O)R¹⁵, -SO₂R¹⁵, -SO₂NR¹⁵R¹⁶, or -NR¹⁵C(=O)R¹⁶, where R¹⁵ and R¹⁶ are independently, hydrogen, optionally substituted lower alkyl, alkenyl, cycloalkyl, optionally substituted heterocycloalkyl,cycloalkyl(lower alkyl), optionally substituted heteroaryl, optionally substituted heteroaryl(lower alkyl) or, together, are -(CH₂)₄₋₆- optionally interrupted by one O, S, NH or N-(C₁₋₂ alkyl) group.

- 56. (previously presented) The compound of claim 55, where R¹³ is optionally substituted lower alkyl, alkenyl, heterocycloalkyl, optionally substituted aryl, optionally substituted aryl(lower alkyl), optionally substituted heteroaryl, optionally substituted heteroaryl(lower alkyl), halo(lower alkyl), -CF₃, halogen, nitro, -CN, -OR¹⁵, -SR¹⁵, -NR¹⁵R¹⁶, -C(=O)R¹⁵, -C(=O)OR¹⁵, -C(=O)OR¹⁵, -C(=O)OR¹⁵, -SO₂OR¹⁵, -SO₂OR¹⁵R¹⁶, or -NR¹⁵C(=O)R¹⁶, where R¹⁵ and R¹⁶ are independently, hydrogen, optionally substituted lower alkyl, alkenyl, cydoalkyl, optionally substituted heterocycloalkyl, cycloalkyl(lower alkyl), optionally substituted aryl, optionally substituted heteroaryl, optionally substituted heteroaryl(lower alkyl) or, together, are -(CH₂)₄₋₆-optionally interrupted by one O, S, NH or N-(C₁₋₂ alkyl) group.
- 57. (previously presented) The compound of claim 51, where R^{14} is independently selected from optionally substituted lower alkyl, optionally substituted aryl, optionally substituted heteroaryl, hydroxy, halogen, $-CF_3$, $-OR^{17}$ $-NR^{17}R^{18}$, $-C(=O)R^{18}$, $-C(=O)OR^{18}$, $-C(=O)NR^{17}R^{18}$, where R^{17} and R^{18} are, independently, hydrogen, lower alkyl, alkenyl, or optionally substituted aryl.
- 58. (previously presented) The compound of claim 56, where n is an integer of 1 to 2.
- 59. (previously presented) The compound of claim 58, where n is 1.
- 60. (previously presented) The compound of claim 51, where R¹ is lower alkyl.
- 61. (previously presented) The compound of claim 51, where R² and R³ are independently selected from hydrogen, lower alkyl, halogen, OR⁹, -NR⁹R¹⁰, where R⁹ and R¹⁰ are independently lower alkyl, substituted lower alkyl, or substituted aryl, or R⁹ and R¹⁰ together are -(CH₂)₄₋₆-

- optionally interrupted by one O, S, NH, N-(aryl), N-(aryl(lower alkyl)), N-(carboxy(lower alkyl)) or N-(optionally substituted C₁₋₂ alkyl) group.
- The compound of claim 56, where R¹³ is independently selected 62. (previously presented) from halogen, optionally substituted aryl, -CF₃, -CH₃, -CN, -OR¹⁵, -C(=O)R¹⁵, -C(=O)OR¹⁵, -C(=O)NR¹⁵R¹⁶, or -CO₂H.
- The compound of claim 51, where R¹⁴ is independently selected 63. (previously presented) from halogen, optionally substituted lower alkyl, -CF₃, -OR¹⁷, aryl, heteroaryl, -NR¹⁷R¹⁸, $-C(=O)R^{17}$, $-C(=O)OR^{17}$, $-C(=O)NR^{17}R^{18}$, or $-CO_2H$, where R^{17} and R^{18} are, independently, lower alkyl, substituted lower alkyl, or substituted aryl, or, together, are -(CH₂)₄₋₆- optionally interrupted by one O, S, NH or N-(C₁₋₂ alkyl) group.
- The compound of claim 51, where R² is 4-methylpiperazinyl, R¹³ is 64. (previously presented) 3-CF₃, and R¹⁴ is 4-F.
- 65. (previously presented) A pharmaceutical composition comprising:
- (a) a therapeutically effective amount of a compound of claim 40; and
- **(b)** a pharmaceutically acceptable excipient.
- 66. (canceled)
- 67. (currently amended) A method of treating an allergic, inflammatory, or autoimmune disorder or disease selected from the group consisting of asthma, atherosclerosis, glomerulonephritis, pancreatitis, restenosis, rheumatoid arthritis, diabetic nephropathy, pulmonary fibrosis, inflammatory bowel disease, Crohn's disease, transplant rejection, and multiple sclerosis, comprising administering a therapeutically effective dose of at least one compound of claim 40 to a mammal in need of such treatment.
- 68. (canceled)
- 69. (canceled)
- 70. (canceled)
- 71. (canceled)